

selected from H, C₁₋₆-alk(en/yn)yl, hydroxy-C₁₋₆-alk(en/yn)yl, C₁₋₆-alk(en/yn)ylsulfanyl-C₁₋₆-alk(en/yn)yl or C₃₋₈-cycloalk(en)yl; or

R⁸ and R⁹ together with the nitrogen form a saturated 3-7 membered heterocyclic ring,
5 and R⁶ is selected from H, C₁₋₆-alk(en/yn)yl, C₁₋₆-alk(en/yn)ylloxy, C₁₋₆-alk(en/yn)ylsulfanyl or C₃₋₈-cycloalk(en)yl, provided that when R⁶ is selected from C₁₋₆-alk(en/yn)ylloxy or C₁₋₆-alk(en/yn)ylsulfanyl then X is CR¹¹R¹², wherein R¹¹ and R¹² independently are selected from H or C₁₋₆ alkyl, and R⁷ is selected from H, C₁₋₆-alk(en/yn)yl or C₃₋₈-cycloalk(en)yl, and R⁹ is selected from H, C₁₋₆-alk(en/yn)yl,
10 hydroxy-C₁₋₆-alk(en/yn)yl, C₁₋₆ alk(en/yn)ylsulfanyl-C₁₋₆-alk(en/yn)yl or C₃₋₈-cycloalk(en)yl;

R¹⁰ is H, C₁₋₆-alk(en/yn)yl, aryl, aryl-C₁₋₆-alk(en/yn)yl, wherein aryl is optionally substituted with a halogen, CF₃, OCF₃, CN, NO₂ or C₁₋₆-alk(en/yn)yl; or an alkali
15 metal;
or a salt thereof, such as a pharmaceutically acceptable salt.

2. The compound of claim 1 wherein X is selected from O or CH₂.

20 3. The compound of any one of claims 1-2 wherein Y is O.

4. The compound of any one of claims 1-2 wherein Y is S.

5. The compound of any one of the preceding claims wherein R¹ is selected from
25 hydrogen, C₁₋₆-alkyl, halogen, phenyl, or phenyl substituted with one or two substituents selected from C₁₋₆-alkyl or C₁₋₆-alkoxy.

6. The compound of any one of the preceding claims wherein R² is selected from
hydrogen; cyano; C₁₋₆-alkyl; halogen; phenyl; phenyl substituted with one or two
30 substituents selected from cyano, C₁₋₆-alkyl, C₁₋₆-alkoxy, or C₁₋₆-alkylsulfonyl; –NR¹³R¹⁴ wherein R¹³ and R¹⁴ together with the nitrogen form a 3-7-membered heterocyclic ring which optionally contains one further heteroatom selected from O, S

or N, such as morpholinyl, or piperidinyl; or monocyclic heteroaryl, such as pyrimidinyl.

7. The compound of any one of the preceeding claims wherein R³ is selected from hydrogen; C₁₋₆-alkyl; halogen; phenyl; phenyl substituted with one or two substituents selected from cyano, C₁₋₆-alkyl, or C₁₋₆-alkoxy; or monocyclic heteroaryl, such as thiophenyl.

8. The compound of any one of the preceeding claims wherein R⁴ is selected from hydrogen, C₁₋₆-alkyl, halogen, phenyl or phenyl substituted with one or two substituents selected from C₁₋₆-alkyl or C₁₋₆-alkoxy.

9. The compound of any one of the preceeding claims wherein R⁵ is phenyl, optionally substituted with a halogen, C₁₋₆-alkyl, C₁₋₆-alkyloxy, C₁₋₆-alkylsulfanyl, halo-C₁₋₆-alkyl.

10. The compound of any one of the preceeding claims wherein R⁶ is selected from H or C₁₋₆-alkyl.

11. The compound of any one of the preceeding claims wherein R⁷ is selected from H or C₁₋₆-alkyl.

12. The compound of any one of the preceding claims wherein R⁸ is selected from H, C₁₋₆-alkyl or C₃₋₈-cycloalkyl.

13. The compound of any one of the preceding claims wherein R⁹ and R^{9'} are independently selected from H or C₁₋₆-alkyl.

14. The compound of any one of the preceding claims wherein R¹⁰ is H.

15. The compound of any one of claims 1-9 or 14 wherein R⁶ and R⁸ together with the nitrogen form a 1-pyrrolidinyl, 1-piperidinyl or 1-azepinyl, optionally substituted with a C₁₋₆-alkyl, and R⁷ is selected from H, C₁₋₆-alk(en/yn)yl or C₃₋₈-cycloalk(en)yl, and

R⁹ and R^{9'} are independently selected from H, C₁₋₆-alk(en/yn)yl, hydroxy-C₁₋₆-alk(en/yn)yl, C₁₋₆ alk(en/yn)ylsulfanyl-C₁₋₆-alk(en/yn)yl or C₃₋₈-cycloalk(en)yl.

16. The compound of any one of claims 1-9 or 14 wherein R⁷ and R⁸ together with the nitrogen form a 1-pyrrolidinyl, 1-piperidinyl or 1-azepinyl, optionally substituted with a C₁₋₆-alkyl, and R⁶ is selected from H, C₁₋₆-alk(en/yn)yl, C₁₋₆-alk(en/yn)ylloxy, C₁₋₆-alk(en/yn)ylsulfanyl or C₃₋₈-cycloalk(en)yl, provided that when R⁶ is selected from C₁₋₆-alk(en/yn)ylloxy or C₁₋₆-alk(en/yn)ylsulfanyl then X is CR¹¹R¹², wherein R¹¹ and R¹² independently are selected from H or C₁₋₆ alkyl, and R⁹ and R^{9'} are independently selected from H, C₁₋₆-alk(en/yn)yl, hydroxy-C₁₋₆-alk(en/yn)yl, C₁₋₆ alk(en/yn)ylsulfanyl-C₁₋₆-alk(en/yn)yl or C₃₋₈-cycloalk(en)yl.

17. The compound of any one of claims 1-9 or 14 wherein R⁸ and R⁹ together with the nitrogen form a 1-pyrrolidinyl, 1-piperidinyl or 1-azepinyl, optionally substituted with a C₁₋₆-alkyl, and R⁶ is selected from H, C₁₋₆-alk(en/yn)yl, C₁₋₆-alk(en/yn)ylloxy, C₁₋₆-alk(en/yn)ylsulfanyl or C₃₋₈-cycloalk(en)yl, provided that when R⁶ is selected from C₁₋₆-alk(en/yn)ylloxy or C₁₋₆-alk(en/yn)ylsulfanyl then X is CR¹¹R¹², wherein R¹¹ and R¹² independently are selected from H or C₁₋₆ alkyl, and R⁷ is selected from H, C₁₋₆-alk(en/yn)yl or C₃₋₈-cycloalk(en)yl, and R^{9'} is selected from H, C₁₋₆-alk(en/yn)yl, hydroxy-C₁₋₆-alk(en/yn)yl, C₁₋₆ alk(en/yn)ylsulfanyl-C₁₋₆-alk(en/yn)yl or C₃₋₈-cycloalk(en)yl.

18. The compound of claim 1 selected from
(S)-1-{2-[2-(4-Fluoro-phenylsulfanyl)-phenoxy]-ethyl}-pyrrolidine-2-carboxylic acid,
(S)-1-{2-[2-(4-tert-Butyl-phenylsulfanyl)-phenoxy]-ethyl}-pyrrolidine-2-carboxylic acid,
(S)-1-{2-[2-(4-Trifluoromethyl-phenylsulfanyl)-phenoxy]-ethyl}-pyrrolidine-2-carboxylic acid,
(S)-1-{2-[2-(3-Fluoro-phenylsulfanyl)-phenoxy]-ethyl}-pyrrolidine-2-carboxylic acid,
(S)-{2-[2-(4-Chloro-phenylsulfanyl)-phenoxy]-ethyl}-pyrrolidine-2-carboxylic acid,
(S)-1-{2-[2-(3-Chloro-phenylsulfanyl)-phenoxy]-ethyl}-pyrrolidine-2-carboxylic acid,

- (S)-1-{2-[2-(3,4-Dichloro-phenylsulfanyl)-phenoxy]-ethyl}-pyrrolidine-2-carboxylic acid,
- (S)-1-{2-[2-(3-Chloro-4-fluoro-phenylsulfanyl)-phenoxy]-ethyl}-pyrrolidine-2-carboxylic acid,
- 5 (S)-1-{2-[2-(3-Chloro-phenoxy)-phenoxy]-ethyl}-pyrrolidine-2-carboxylic acid,
- (S)-1-{2-[2-(4-Chloro-phenoxy)-phenoxy]-ethyl}-pyrrolidine-2-carboxylic acid,
- (S)-1-{2-[2-(4-Methoxy-phenoxy)-phenoxy]-ethyl}-pyrrolidine-2-carboxylic acid,
- (S)-1-{2-[2-(3,4-Difluoro-phenoxy)-phenoxy]-ethyl}-pyrrolidine-2-carboxylic acid,
- 1-{2(R/S)-[2-(4-Chloro-phenoxy)-phenoxy]-propyl}-pyrrolidine-2(S)-carboxylic acid,
- 10 1-{2(R/S)-[2-(3,4-Difluoro-phenoxy)-phenoxy]-propyl}-pyrrolidine-2(S)-carboxylic acid,
- (S)-1-{2-[2-(3-Fluoro-phenoxy)-phenoxy]-ethyl}-pyrrolidine-2-carboxylic acid,
- 1-{2(R/S)-[2-(3-Fluoro-phenoxy)-phenoxy]-propyl}-pyrrolidine-2(S)-carboxylic acid,
- 1-{2(R/S)-[2-(3-Fluoro-phenylsulfanyl)-phenoxy]-propyl}-pyrrolidine-2(S)-
- 15 carboxylic acid,
- 1-{2(R/S)-[2-(3-Chloro-phenylsulfanyl)-phenoxy]-propyl}-pyrrolidine-2(S)-carboxylic acid,
- ({2-[2-(4-tert-Butyl-phenylsulfanyl)-phenoxy]-ethyl}-N-ethyl-amino)-acetic acid,
- 2-{3-[2-(4-tert-Butyl-phenylsulfanyl)-phenoxy]-pyrrolidin-1-yl}-propionic acid,
- 20 ({2-[2-(3-Chloro-phenylsulfanyl)-phenoxy]-ethyl}-N-methyl-amino)-acetic acid,
- ({2-[2-(3-Chloro-4-fluoro-phenylsulfanyl)-phenoxy]-ethyl}-N-methyl-amino)-acetic acid,
- {2-[2-(4-tert-Butyl-phenylsulfanyl)-phenoxy-methyl]-piperidin-1-yl}-acetic acid,
- ({2-[2-(3-Fluoro-phenylsulfanyl)-phenoxy]-ethyl}-N-methyl-amino)-acetic acid,
- 25 {4-[2-(4-tert-Butyl-phenylsulfanyl)-phenoxy]-piperidin-1-yl}-acetic acid,
- (N-2-propyl-{2-[2-(4-trifluoromethyl-phenylsulfanyl)-phenoxy]-ethyl}-amino)-acetic acid,
- ({2-[2-(3,4-Dichloro-phenylsulfanyl)-phenoxy]-ethyl}-N-ethyl-amino)-acetic acid,
- (N-Ethyl-{2-[2-(4-methylsulfanyl-phenylsulfanyl)-phenoxy]-ethyl}-amino)-acetic
- 30 acid,
- 2-{3-[2-(3,4-Dichloro-phenylsulfanyl)-phenoxy]-pyrrolidin-1-yl}-propionic acid,
- (S)-{3-[2-(4-tert-Butyl-phenylsulfanyl)-phenoxy]-pyrrolidin-1-yl}-acetic acid,

- ({2-[2-(3-Chloro-4-fluoro-phenylsulfanyl)-phenoxy]-ethyl}-N-ethyl-amino)-acetic acid,
(N-2-propyl-{2-[2-(4-methylsulfanyl-phenylsulfanyl)-phenoxy]-ethyl}-amino)-acetic acid,
5 {3-[2-(4-tert-Butyl-phenylsulfanyl)-phenoxy]-pyrrolidin-1-yl}-acetic acid,
({2-[2-(3-Chloro-phenylsulfanyl)-phenoxy]-ethyl}-N-ethyl-amino)-acetic acid,
({2-[2-(4-Chloro-phenylsulfanyl)-phenoxy]-ethyl}-N-methyl-amino)-acetic acid,
{4-[2-(3,4-Dichloro-phenylsulfanyl)-phenoxy]-piperidin-1-yl}-acetic acid,
2-{3-[2-(4-Trifluoromethyl-phenylsulfanyl)-phenoxy]-pyrrolidin-1-yl}-propionic
10 acid,
({2-[2-(4-tert-Butyl-phenylsulfanyl)-phenoxy]-ethyl}-N-2-propyl-amino)-acetic acid
({2-[2-(4-tert-Butyl-phenylsulfanyl)-phenoxy]-ethyl}-N-methyl-amino)-acetic acid,
{2-[2-(4-Methylsulfanyl-phenylsulfanyl)-phenoxy-methyl]-piperidin-1-yl}-acetic acid,
({2-[2-(3,4-Dichloro-phenylsulfanyl)-phenoxy]-ethyl}-N-methyl-amino)-acetic acid,
15 (N-Methyl-{2-[2-(4-trifluoromethyl-phenylsulfanyl)-phenoxy]-ethyl}-amino)-acetic acid,
2-{3(R)-[2-(4-tert-Butyl-phenylsulfanyl)-phenoxy]-pyrrolidin-1-yl}-propionic acid,
2-{3(R)-[2-(3,4-Dichloro-phenylsulfanyl)-phenoxy]-pyrrolidin-1-yl}-propionic acid,
2-[3(R)-(2-(4-methylphenyl)-sulfanyl-phenoxy)-pyrrolidin-1-yl]-propionic acid,
20 {3(R)-[2-(4-tert-Butyl-phenylsulfanyl)-phenoxy]-pyrrolidin-1-yl}-acetic acid,
2-{3(R)-[2-(4-Trifluoromethyl-phenylsulfanyl)-phenoxy]-pyrrolidin-1-yl}-propionic acid,
2-{3(R)-[2-(4-Chloro-phenylsulfanyl)-phenoxy]-pyrrolidin-1-yl}-propionic acid,
({1-[2-(3-Chloro-phenylsulfanyl)-phenoxy-methyl]-propyl}-N-ethyl-amino)-acetic
25 acid,
({1-[2-(3,4-Dichloro-phenylsulfanyl)-phenoxy]-butan-2-yl}-N-ethyl-amino)-acetic acid,
({1-[2-(3,4-Dichloro-phenylsulfanyl)-phenoxy]-butan-3-methyl-2-yl}-N-ethyl-amino)-acetic acid,
30 ({1-[2-(3-Chloro-4-fluoro-phenylsulfanyl)-phenoxy]-butan-2-yl}-N-ethyl-amino)-acetic acid,
({1-[1-(3-Chloro-phenylsulfanyl)-phenoxy]-propan-2-yl}-N-ethyl-amino)-acetic acid,

({1-[2-(3-Chloro-4-fluoro-phenylsulfanyl)-phenoxy]-butan-4-methyl-2-yl})-N-ethyl-amino)-acetic acid,

({1-[2-(3-Chloro-4-fluoro-phenylsulfanyl)-phenoxy]propan-2-yl}-N-ethyl-amino)-acetic acid,

5 (S)-{1-[2-(3-Chloro-phenylsulfanyl)-phenoxy]-propan-2-yl}-N-methyl-amino)-acetic acid,

(S)-({1-[2-(3-Chloro-phenylsulfanyl)-phenoxy]-propan-2-yl}-N-ethyl-amino)-acetic acid,

10 ({1-[2-(3,4-Dichloro-phenylsulfanyl)-phenoxy]-propan-2-yl}-N-ethyl-amino)-acetic acid,

({1-[2-(4-Chloro-phenylsulfanyl)-phenoxy]-propan-2-yl}-N-ethyl-amino)-acetic acid,

({1-[2-(3-Chloro-phenylsulfanyl)-phenoxy]methyl]-propyl}-N-methyl-amino)-acetic acid,

15 ({1-[2-(4-Chloro-phenylsulfanyl)-phenoxy]methyl]-propyl}-N-ethyl-amino)-acetic acid,

(N-Ethyl-{1-[2-(3-fluoro-phenylsulfanyl)-phenoxy]methyl]-propyl}-amino)-acetic acid,

(R)-({2-[2-(3,4-Dichloro-phenylsulfanyl)-phenoxy]-1-methyl-ethyl}-N-ethyl-amino)-acetic acid,

20 (S)-(2-{2-[2-(4-Chloro-phenoxy)-phenoxy]-propyl}-N-methyl-amino)-acetic acid,

(R)-(2-{2-[2-(3-Chloro-phenylsulfanyl)-phenoxy]-}-propyl)-N-methyl-amino)-acetic acid,

({2-[2-(3-Fluoro-phenylsulfanyl)-phenoxy]-propyl}-N-methyl-amino)-acetic acid,

({2-[2-(3-Chloro-phenylsulfanyl)-phenoxy]-propan-1-yl}-N-ethyl-amino)-acetic acid,

25 ({1-[2-(3-Chloro-phenylsulfanyl)-phenoxy]-3-methyl-butan-2-yl}-N-methyl-amino)-acetic acid,

({3-methyl-1-[2-(4-trifluoromethyl-phenylsulfanyl)-phenoxy]-butan-2-yl}-N-ethyl-amino)-acetic acid,

30 ({1-[2-(3-Chloro-4-fluoro-phenylsulfanyl)-phenoxy]-butan-2-yl}-N-methyl-amino)-acetic acid,

(S)-(1-{2-[2-(3-Chloro-4-fluoro-phenylsulfanyl)-phenoxy]-propan-2-yl}-N-methyl-amino)-acetic acid,

- (S)-(2-{2-[2-(3-Fluoro-phenylsulfanyl)-phenoxy]-propyl}-N-methyl-amino)-acetic acid,
- ((1-[2-(4-tert-Butyl-phenylsulfanyl)-phenoxy]-3-methyl-butan-2-yl)-N-ethyl-amino)-acetic acid,
- 5 (S)-({1-[2-(3,4-Dichloro-phenylsulfanyl)-phenoxy]-propan-2-yl}-N-methyl-amino)-acetic acid,
- ((1-[2-(3-Chloro-4-fluoro-phenylsulfanyl)-phenoxy]-3-methyl-butan-2-yl)-N-methyl-amino)-acetic acid,
- ((1-[2-(4-tert-Butyl-phenylsulfanyl)-phenoxy]-3-methyl-propan-2-yl)-N-ethyl-
- 10 amino)-acetic acid,
- ((2-[2-(3-Chloro-4-fluoro-phenylsulfanyl)-phenoxy]-propan-1-yl)-N-ethyl-amino)-acetic acid,
- ((2-[2-(4-methoxy-phenylsulfanyl)-phenoxy]-propan-1-yl)-N-Cyclohexyl-amino)-acetic acid,
- 15 { [2-(2-(4-methylsulfanyl)-phenoxy)-propan-1-yl]-N-cyclohexyl-amino}-acetic acid,
- ((2-[2-(3-Chloro-phenylsulfanyl)-phenoxy]-propan-1-yl)-N-cyclohexyl-amino)-acetic acid,
- (S)-1-{3-[2-(3-Fluoro-phenylsulfanyl)-phenyl]-propyl}-pyrrolidine-2-carboxylic acid,
- (S)-2-({2-[3-(3-Fluoro-phenylsulfanyl)-biphenyl-4-yloxy]-ethyl}-methyl-amino)-
- 20 propionic acid,
- ((2-[3-(3-Fluoro-phenylsulfanyl)-biphenyl-4-yloxy]-ethyl)-methyl-amino)-acetic acid,
- (S)-1-{2-[4-Chloro-2-(3-fluoro-phenylsulfanyl)-phenoxy]-ethyl}-pyrrolidine-2-carboxylic acid,
- 25 (S)-1-{2-[3-Chloro-2-(3-fluoro-phenylsulfanyl)-phenoxy]-ethyl}pyrrolidine-2-carboxylic acid,
- (S)-1-{2-[5-Chloro-2-(3-fluoro-phenylsulfanyl)-phenoxy]-ethyl}pyrrolidine-2-carboxylic acid,
- (S)-1-{2-[4-Cyano-2-(3-fluoro-phenylsulfanyl)-phenoxy]-ethyl}-pyrrolidine-2-
- 30 carboxylic acid
- (S)-1-[2-(5-Chloro-2-phenylsulfanyl)-phenoxy]-ethyl]pyrrolidine-2-carboxylic acid,
- (S)-1-{2-[3-(3-Fluoro-phenylsulfanyl)-biphenyl-4-yloxy]-ethyl}-pyrrolidine-2-carboxylic acid,

(S)-{2-[4'-Methoxy-3-(3-fluoro-phenylsulfanyl)-biphenyl-4-yloxy]-ethyl}-pyrrolidine-2-carboxylic acid,

(S)-{2-[4'-Cyano-3-(3-fluoro-phenylsulfanyl)-biphenyl-4-yloxy]-ethyl}-pyrrolidine-2-carboxylic acid,

5 (S)-1-{2-[4'-Cyano-4-(3-fluoro-phenylsulfanyl)-biphenyl-3-yloxy]-ethyl}-pyrrolidine-2-carboxylic acid,

(S)-1-{2-[2-(3-Fluoro-phenylsulfanyl)-5-thiophen-3-yl-phenoxy]-ethyl}-pyrrolidine-2-carboxylic acid,

10 (S)-1-{2-[2-(3-Fluoro-phenylsulfanyl)-4-pyrimidin-5-yl-phenoxy]-ethyl}-pyrrolidine-2-carboxylic acid,

(S)-1-{2-[3-(3-Fluoro-phenylsulfanyl)-3-methanesulfonyl-biphenyl-4-yloxy]-ethyl}-pyrrolidine-2(S)-carboxylic acid,

(S)-1-{2-[2-(3-Fluoro-phenylsulfanyl)-4-morpholin-4-yl-phenoxy]-ethyl}-pyrrolidine-2-carboxylic acid,

15 (S)-1-{2-[2-(3-Fluoro-phenylsulfanyl)-4-piperidin-1-yl-phenoxy]-ethyl}-pyrrolidine-2-carboxylic acid,

or a pharmaceutically acceptable salt thereof.

19. A pharmaceutical composition comprising a compound according to any one of
20 claims 1-18 and a pharmaceutically acceptable carrier or diluent.

20. The use of a compound according to any one of claims 1-18 for the preparation of a medicament for the treatment of post-traumatic stress disorder or a disease selected from the group consisting of schizophrenia, including both the positive and the
25 negative symptoms of schizophrenia and other psychoses, and in the improvement of cognition in conditions where the cognitive processes are diminished, i.e. Alzheimer's disease, multi-infarct dementia, AIDS dementia, Huntington's disease, Parkinson's disease, amyotrophic lateral sclerosis or diseases wherein the brain is damaged by inner or outer influence, such as trauma to the head or stroke, and convulsive
30 disorders such as epilepsy, spasticity or myoclonus.

21. A method for the treatment of a disease or disorder selected from the group consisting of post-traumatic stress disorder, the positive and the negative symptoms of

schizophrenia, including both the positive and the negative symptoms of schizophrenia and other psychoses, and in the improvement of cognition in conditions where the cognitive processes are diminished, i.e. Alzheimer's disease, multi-infarct dementia, AIDS dementia, Huntington's disease, Parkinson's disease, amyotrophic lateral sclerosis or diseases wherein the brain is damaged by inner or outer influence, such as trauma to the head or stroke, and convulsive disorders such as epilepsy, spasticity or myoclonus in a living animal body, including a human, comprising administering to a subject in need thereof a therapeutically effective amount of a compound according to any one of claims 1-18.